

101033,055

(FILE 'HOME' ENTERED AT 09:40:34 ON 25 JUN 2004)

FILE 'REGISTRY' ENTERED AT 09:40:55 ON 25 JUN 2004

L1 1 S (169590-42-5)/RN  
L2 1 S (76-42-6)/RN

FILE 'CAPLUS' ENTERED AT 09:41:40 ON 25 JUN 2004

L3 28 S L1 AND L2  
L4 973 S L1  
L5 732 S L2  
L6 178 S L4 AND PAIN?  
L7 8 S L6 AND SYNERG?  
L8 56 S L4 AND (OPIOID? OR OPIATE? OR MORPHIN?)  
L9 32 S L8 NOT L3  
L10 8 S L1 AND PAIN? AND SYNERG?

FILE 'MEDLINE, SCISEARCH, BIOSIS' ENTERED AT 09:47:31 ON 25 JUN 2004

L11 2083 S L1  
L12 2 S L11 AND PAIN? AND SYNERG?  
L13 89 S L11 AND PAIN? AND (ACETAMINOPHEN? OR IBUPROFEN? OR ASPIRIN?)  
L14 69 DUP REM L13 (20 DUPLICATES REMOVED)

FILE 'USPATFULL' ENTERED AT 09:50:44 ON 25 JUN 2004

L15 301 S L1  
L16 221 S L15 AND (ASPIRIN? OR ACETAMIN? OR IBUPROFEN?)  
L17 169 S L16 AND PAIN?  
L18 58 S L17 AND (PAIN)/CLM  
L19 23 S L18 AND SYNERG?

FILE 'USPATFULL' ENTERED AT 09:54:25 ON 25 JUN 2004

L20 21 S L1 AND L2  
L21 1480 S (COX?)/CLM  
L22 54 S (OPIOID? OR OPIAT? OR MORPHIN? OR OXYCOD?)/CLM AND L21

FILE 'WPIDS' ENTERED AT 09:57:47 ON 25 JUN 2004

L23 0 S L1

FILE 'CAPLUS' ENTERED AT 09:58:04 ON 25 JUN 2004

L24 973 S L1  
L25 178 S L24 AND PAIN?

FILE 'STNGUIDE' ENTERED AT 09:59:42 ON 25 JUN 2004

FILE 'CAPLUS' ENTERED AT 10:01:37 ON 25 JUN 2004

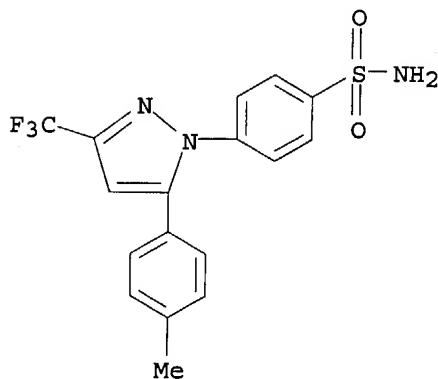
FILE 'STNGUIDE' ENTERED AT 10:03:17 ON 25 JUN 2004

FILE 'MEDLINE, SCISEARCH' ENTERED AT 10:04:01 ON 25 JUN 2004

L26 836 S L1  
L27 155 S L26 AND PAIN?

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L1 ANSWER 1 OF 1 REGISTRY COPYRIGHT 2004 ACS on STN  
 RN 169590-42-5 REGISTRY  
 CN Benzenesulfonamide, 4-[5-(4-methylphenyl)-3-(trifluoromethyl)-1H-pyrazol-1-yl]- (9CI) (CA INDEX NAME)  
 OTHER NAMES:  
 CN 4-[5-(4-Methylphenyl)-3-(trifluoromethyl)-1H-pyrazol-1-yl]benzenesulfonamide  
 CN Celebrex  
 CN Celecoxib  
 CN Celocoxib  
 CN SC 58635  
 CN YM 177  
 FS 3D CONCORD  
 DR 184007-95-2, 194044-54-7  
 MF C17 H14 F3 N3 O2 S  
 CI COM  
 SR US Adopted Names Council (USAN)  
 LC STN Files: ADISINSIGHT, ADISNEWS, AGRICOLA, ANABSTR, BIOBUSINESS, BIOSIS, BIOTECHNO, CA, CANCERLIT, CAPLUS, CASREACT, CBNB, CEN, CHEMCATS, CIN, CSChem, DDFU, DIOGENES, DRUGU, EMBASE, HSDB\*, IMSCoSEARCH, IMSDRUGNEWS, IMSPATENTS, IMSRESEARCH, IPA, MEDLINE, MRCK\*, PHAR, PROMT, PROUSDDR, PS, RTECS\*, SYNTHLINE, TOXCENTER, USAN, USPAT2, USPATFULL  
 (\*File contains numerically searchable property data)  
 DT.CA Caplus document type: Book; Conference; Dissertation; Journal; Patent  
 RL.P Roles from patents: ANST (Analytical study); BIOL (Biological study); PREP (Preparation); PROC (Process); PRP (Properties); RACT (Reactant or reagent); USES (Uses)  
 RLD.P Roles for non-specific derivatives from patents: BIOL (Biological study); PREP (Preparation); PROC (Process); PRP (Properties); USES (Uses)  
 RL.NP Roles from non-patents: ANST (Analytical study); BIOL (Biological study); MSC (Miscellaneous); PREP (Preparation); PROC (Process); PRP (Properties); RACT (Reactant or reagent); USES (Uses)  
 RLD.NP Roles for non-specific derivatives from non-patents: BIOL (Biological study); PREP (Preparation); PROC (Process); PRP (Properties); USES (Uses)



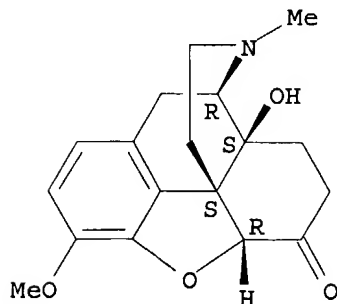
\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

962 REFERENCES IN FILE CA (1907 TO DATE)  
 23 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA  
 973 REFERENCES IN FILE CAPLUS (1907 TO DATE)

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L2 ANSWER 1 OF 1 REGISTRY COPYRIGHT 2004 ACS on STN  
 RN 76-42-6 REGISTRY  
 CN Morphinan-6-one, 4,5-epoxy-14-hydroxy-3-methoxy-17-methyl-, (5 $\alpha$ )-  
 (9CI) (CA INDEX NAME)  
 OTHER CA INDEX NAMES:  
 CN Codeinone, 7,8-dihydro-14-hydroxy- (6CI, 7CI)  
 CN Morphinan-6-one, 4,5 $\alpha$ -epoxy-14-hydroxy-3-methoxy-17-methyl- (8CI)  
 OTHER NAMES:  
 CN (-)-Oxycodone  
 CN 14-Hydroxydihydrocodeinone  
 CN 3-O-(Methyl)oxymorphone  
 CN 6-Oxo-14-hydroxy-7,8-dihydrocodeine  
 CN 7,8-Dihydro-14-hydroxycodeinone  
 CN Dihydro-14-hydroxycodeinone  
 CN Dihydrohydroxycodeinone  
 CN Dihydrone  
 CN NSC 19043  
 CN Oxanest  
 CN Oxicon  
 CN Oxycodeinone  
 CN Oxycodone  
 CN Oxymorphone 3-methyl ether  
 FS STEREOSEARCH  
 MF C18 H21 N O4  
 CI COM  
 LC STN Files: ADISINSIGHT, ADISNEWS, AGRICOLA, ANABSTR, BEILSTEIN\*,  
 BIOBUSINESS, BIOSIS, BIOTECHNO, CA, CANCERLIT, CAOLD, CAPLUS, CASREACT,  
 CBNB, CEN, CHEMCATS, CHEMINFORMRX, CHEMLIST, CIN, CSCHEM, DDFU,  
 DIOGENES, DRUGU, EMBASE, GMELIN\*, HSDB\*, IFICDB, IFIPAT, IFIUDB, IPA,  
 MEDLINE, MRCK\*, MSDS-OHS, NAPRALERT, NIOSHTIC, PHAR, PROMT, PROUSDDR,  
 PS, RTECS\*, SPECINFO, TOXCENTER, USAN, USPAT2, USPATFULL  
 (\*File contains numerically searchable property data)  
 Other Sources: EINECS\*\*, WHO  
 (\*\*Enter CHEMLIST File for up-to-date regulatory information)  
 DT.CA Caplus document type: Conference; Journal; Patent  
 RL.P Roles from patents: ANST (Analytical study); BIOL (Biological study);  
 FORM (Formation, nonpreparative); MSC (Miscellaneous); PREP  
 (Preparation); PROC (Process); PRP (Properties); RACT (Reactant or  
 reagent); USES (Uses); NORL (No role in record)  
 RLD.P Roles for non-specific derivatives from patents: BIOL (Biological  
 study); PREP (Preparation); USES (Uses)  
 RL.NP Roles from non-patents: ANST (Analytical study); BIOL (Biological  
 study); FORM (Formation, nonpreparative); OCCU (Occurrence); PREP  
 (Preparation); PROC (Process); PRP (Properties); RACT (Reactant or  
 reagent); USES (Uses); NORL (No role in record)  
 RLD.NP Roles for non-specific derivatives from non-patents: ANST (Analytical  
 study); BIOL (Biological study)

Absolute stereochemistry.



\*\*PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT\*\*

728 REFERENCES IN FILE CA (1907 TO DATE)  
15 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA  
732 REFERENCES IN FILE CAPLUS (1907 TO DATE)  
32 REFERENCES IN FILE CAOLD (PRIOR TO 1967)

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L25 ANSWER 178 OF 178 CAPLUS COPYRIGHT 2004 ACS on STN  
AN 1997:231026 CAPLUS  
DN 126:264035

TI Synthesis and Biological Evaluation of the 1,5-Diarylpyrazole Class of  
Cyclooxygenase-2 Inhibitors: Identification of 4-[5-(4-Methylphenyl)-3-  
(trifluoromethyl)-1H-pyrazol-1-yl]benzenesulfonamide (SC-58635, Celecoxib)

AU Penning, Thomas D.; Talley, John J.; Bertenshaw, Stephen R.; Carter,  
Jeffery S.; Collins, Paul W.; Docter, Stephen; Graneto, Matthew J.; Lee,  
Len F.; Malecha, James W.; Miyashiro, Julie M.; Rogers, Roland S.; Rogier,  
D. J.; Yu, Stella S.; Anderson, Gary D.; Burton, Earl G.; Cogburn, J.  
Nita; Gregory, Susan A.; Koboldt, Carol M.; Perkins, William E.; Seibert,  
Karen; Veenhuizen, Amy W.; Zhang, Yan Y.; Isakson, Peter C.

CS Departments of Chemistry Inflammatory Diseases Research and Molecular  
Pharmacology, Searle Research and Development, Skokie, IL, 60077, USA

SO Journal of Medicinal Chemistry (1997), 40(9), 1347-1365

CODEN: JMCMAR; ISSN: 0022-2623

PB American Chemical Society

DT Journal

LA English

AB Sulfonamide-containing 1,5-diarylpyrazole derivs. were prepared and evaluated  
for their ability to block cyclooxygenase-2 (COX-2) in vitro and in vivo.  
Extensive structure-activity relationship work was carried out within this  
series, and a number of potent and selective inhibitors of COX-2 were  
identified. Since an early structural lead exhibited an unacceptably long  
plasma half-life, a number of pyrazole analogs containing potential metabolic  
sites were evaluated further in vivo in an effort to identify compds. with  
acceptable pharmacokinetic profiles. This work led to the identification  
of SC-58635 (celecoxib, I), which is currently in phase III clin. trials  
for the treatment of rheumatoid arthritis and osteoarthritis.

IT Pain

(hyperalgesia; diarylpyrazoles as cyclooxygenase 2 inhibitors)

IT 169590-42-5P 170569-50-3P 170569-69-4P 170569-75-2P  
170569-83-2P 170569-85-4P 170569-88-7P 170569-91-2P 170570-25-9P  
170570-80-6P 170570-81-7P 170571-00-3P 170571-05-8P 170571-29-6P  
170571-71-8P 170571-92-3P 170571-97-8P 170572-00-6P 170572-05-1P  
170572-08-4P 188816-97-9P 188816-98-0P 188816-99-1P 188817-00-7P  
188817-07-4P

RL: BAC (Biological activity or effector, except adverse); BSU (Biological  
study, unclassified); RCT (Reactant); SPN (Synthetic preparation); BIOL  
(Biological study); PREP (Preparation); RACT (Reactant or reagent)  
(diarylpyrazoles as cyclooxygenase 2 inhibitors)

L25 ANSWER 176 OF 178 CAPLUS COPYRIGHT 2004 ACS on STN  
AN 1997:521520 CAPLUS  
DN 127:171270  
TI Outcome of specific COX-2 inhibition in rheumatoid arthritis  
AU Lipsky, Peter E.; Isakson, Peter C.  
CS Rheumatic Diseases Division, Department of Internal Medicine, University  
Texas Southwestern Medical Center at Dallas, Dallas, TX, 75235-8884, USA  
SO Journal of Rheumatology, Supplement (1997), 49(Progress toward a New Class  
of Therapeutics: Selective COX-2 Inhibition), 9-14  
CODEN: JRSUDX; ISSN: 0380-0903  
PB Journal of Rheumatology  
DT Journal  
LA English

=> d 176 ab

L25 ANSWER 176 OF 178 CAPLUS COPYRIGHT 2004 ACS on STN  
AB We reviewed data suggesting the hypothesis that specific inhibition of the  
inducible isoform of cyclooxygenase, COX-2, would provide therapeutic  
benefit in patients with rheumatoid arthritis (RA) with less  
gastrointestinal toxicity and presented the results of a therapeutic trial  
to test this hypothesis. Various doses of the selective COX-2 inhibitor,  
celecoxib, or placebo were used to treat patients with RA in a 4 wk,  
double blind, placebo controlled trial. Celecoxib provided significant  
improvement in patient global assessment, morning stiffness, and the number  
of **painful** and tender joints compared with placebo. In addition,  
the number of withdrawals in celecoxib treated patients was significantly  
less than in the placebo group. No significant adverse events and no  
difference in the total number of adverse events were noted between the  
placebo and celecoxib groups. At the doses employed, celecoxib inhibited  
only COX-2 and not COX-1. Specific COX-2 inhibition with celecoxib causes  
significant improvement in the signs and symptoms of RA.

L27 ANSWER 155 OF 155 MEDLINE on STN  
AN 97393128 MEDLINE  
DN PubMed ID: 9249645  
TI Outcome of specific COX-2 inhibition in rheumatoid arthritis.  
AU Lipsky P E; Isakson P C  
CS Department of Internal Medicine, University of Texas Southwestern Medical  
Center, Dallas 75235-8884, USA.  
SO Journal of rheumatology, (1997 Jul) 24 Suppl 49 9-14. Ref: 21  
Journal code: 7501984. ISSN: 0315-162X.  
CY Canada  
DT (CLINICAL TRIAL)  
(CONTROLLED CLINICAL TRIAL)  
Journal; Article; (JOURNAL ARTICLE)  
General Review; (REVIEW)  
(REVIEW, TUTORIAL)  
LA English  
FS Priority Journals  
EM 199709  
ED Entered STN: 19970916  
Last Updated on STN: 20000303  
Entered Medline: 19970904

L27 ANSWER 154 OF 155 MEDLINE on STN  
AN 97393130 MEDLINE  
DN PubMed ID: 9249647  
TI Pain management in osteoarthritis: the role of COX-2 inhibitors.  
AU Lane N E  
CS Department of Medicine, University of California at San Francisco 94143,  
USA.. nelane@itsa.ucsf.edu  
NC AG05407 (NIA)  
AR20684 (NIAMS)  
SO Journal of rheumatology, (1997 Jul) 24 Suppl 49 20-4. Ref: 30  
Journal code: 7501984. ISSN: 0315-162X.  
CY Canada  
DT Journal; Article; (JOURNAL ARTICLE)  
General Review; (REVIEW)  
(REVIEW, TUTORIAL)  
LA English  
FS Priority Journals  
EM 199709  
ED Entered STN: 19970916  
Last Updated on STN: 20000303  
Entered Medline: 19970904

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